



**Manchester  
Metropolitan  
University**

---

Borillo, GC, Tadano, YS, Godoi, AFL, Santana, SSM, Weronka, FM, Penteadó Neto, RA, Rempel, D, Yamamoto, CI, Potgieter, SS, Potgieter, JH and Godoi, RHM (2015) Effectiveness of Selective Catalytic Reduction (SCR) systems on reducing gaseous emissions from an engine using Diesel and Biodiesel Blends. *Environmental Science and Technology*, 49 (5). pp. 3246-3251. ISSN 0013-936X

---

**Downloaded from:** <https://e-space.mmu.ac.uk/620398/>

**Publisher:** American Chemical Society

**DOI:** <https://doi.org/10.1021/es505701r>

Please cite the published version

<https://e-space.mmu.ac.uk>

# **Effectiveness of Selective Catalytic Reduction (SCR) systems on reducing gaseous emissions from an engine using Diesel and Biodiesel Blends**

Guilherme C. Borillo<sup>1</sup>, Yara S. Tadano<sup>2</sup> Ana Flavia L. Godoi<sup>1</sup>; Simone S. M. Santana<sup>1</sup>; Fernando M. Weronca<sup>1</sup>; Renato A. Penteado Neto<sup>3</sup>; Dennis Rempel<sup>3</sup>; Carlos I. Yamamoto<sup>4</sup>; Sanja Potgieter-Verma<sup>5</sup>; J. Herman Potgieter<sup>6</sup>; Ricardo H. M. Godoi<sup>1\*</sup>

\*Corresponding Author: Environmental Engineering Department, Federal University of Parana, 210 Francisco H. dos Santos St., Curitiba, PR, 81531-980 Brazil. Tel.: +55 41 3361-3482. E-mail address: rhmgodoi@ufpr.br (R. H. M. Godoi).

1- Environmental Engineering Department, Federal University of Parana, Curitiba, PR, Brazil.

2- Mathematics Department, Federal University of Technology - Parana, Ponta Grossa, PR, Brazil.

3- Vehicle Emissions Laboratory, Lactec Institute, Curitiba, PR, Brazil.

4- Chemical Engineering Department, Federal University of Parana, Curitiba, PR, Brazil.

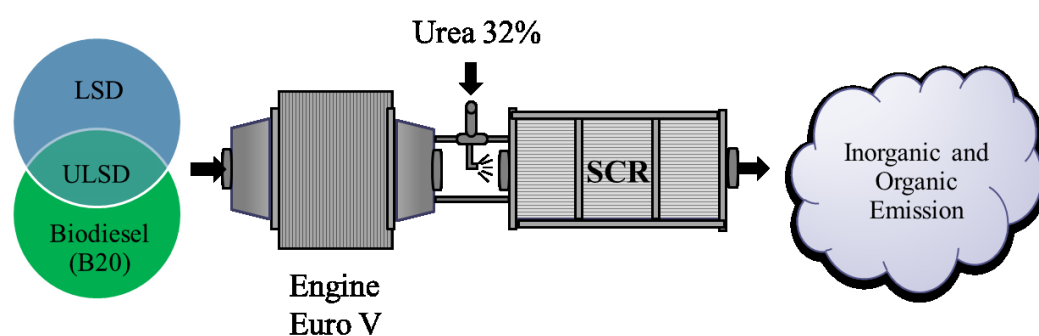
5- Division of Chemistry and Environmental Science, School of Science and the Environment, Manchester Metropolitan University, Manchester, UK

6- School of Chemical and Metallurgical Engineering, University of the Witwatersrand, Johannesburg, South Africa.

ABSTRACT

There is an urgent and pressing need to further understand petroleum-based emission control systems. To date, a limited number of emission studies have reported on the effects on automotive emissions when vehicles equipped with Selective Catalytic Reduction (SCR) systems run on a mixture of regular petroleum-based and biodiesel. The aim of this investigation was to quantify organic and inorganic gas emissions from a four-cylinder diesel engine equipped with urea-SCR system. Using a bench dynamometer, the emissions from the following mixtures were evaluated using an FTIR spectrometer: low sulphur diesel (LSD), ultra-low sulphur diesel (ULSD) and a blend of 20 % soybean biodiesel and 80% ULSD (B20). Our results confirmed that the use of the SCR system yields statistically significant ( $p < 0.05$ ) lower  $\text{NO}_x$  emissions in comparison to all the studied fuels. The LSD and ULSD fuels also significantly reduced emissions of compounds with high photochemical ozone creation potential, such as formaldehyde. However, the SCR system produced significantly ( $p < 0.05$ ) higher emissions of  $\text{N}_2\text{O}$  comparing the used fuels. In the case of LSD, the  $\text{NH}_3$  emissions were elevated and in the case of ULSD and B20 fuels, the non-methane hydrocarbon (NMHC) and total hydrocarbon (HCD) emissions were significantly higher.

**Keywords:** Selective Catalytic Reduction (SCR); biodiesel; hydrocarbons; diesel; emissions; gaseous pollutants.



## 1. Introduction

There is an urgent and pressing need for in-depth understanding of petroleum-based emission control systems. Global pressure to meet emission standards lead to the

development and use of new engine technologies and as of late also for the use of new fuels and fuel blends, such as ultra-low sulphur diesel and biodiesel blends.

Emissions depend on a variety of factors, such as engine technology, maintenance and emission control technology,<sup>1</sup> as well as the type and quality of the employed fuel. Besides the greenhouse gas pollutants with global warming potential, it is widely known that engine exhaust systems produce also organic gases that have an impact on photochemical ozone and other secondary pollutants' formation. Among such different gases emitted by petroleum-based systems, nitrogen oxides (NO<sub>x</sub>) are one of the major threats to the environment and therefore its emission in diesel engines has been widely investigated.<sup>2-5</sup> NO<sub>x</sub> suppression strategies consist of combustion controls, such as Selective Catalytic Reduction (SCR) systems, using a urea solution as reducing agent, a well-established technique of stationary diesel engines.<sup>6-8</sup> Biodiesel seems to be a promising alternative, as it can be used in diesel engines without major modifications,<sup>9</sup> reducing qualitative and quantitatively several pollutant emissions.<sup>10-14</sup> The use of biofuels and fuel blends, in combination with exhaust aftertreatment systems as a means of mitigating emissions, are promising and therefore the topic of this investigation.

New standard guidelines are being established worldwide concerning heavy-duty diesel engine emissions, aiming mostly at the simultaneous reduction of particles and NO<sub>x</sub> (Euro V and Euro VI regulations in Europe and 40 Code of Federal Regulations 86.007-11).<sup>15</sup> In Brazil, the ruling legislation is equivalent to the Euro V emission standards and it was established on January 1<sup>st</sup>, 2012, as a result of the seventh stage of the Program to Control Vehicular Air Pollution (PROCONVE, in Portuguese). In order to achieve the Brazilian air quality guidelines, the sulphur content of diesel fuels was reduced and new aftertreatment systems have been implemented, with the urea-SCR (Selective Catalytic Reduction) system being mostly utilized.<sup>4,5,16</sup>

To date, a limited number of emission studies have reported on the effects of biodiesel additions to regular petroleum-based diesel on emissions from vehicles equipped with Selective Catalytic Reduction (SCR) systems.

In order to fill the gap, the aim of this investigation was to quantify organic and inorganic gas emissions (gas- and particle-phase) from a four-cylinder diesel engine equipped with an urea-SCR system using Diesel or Biodiesel blends.

## 2. Materials and methods

In this study, we used an engine dynamometer following the European Steady Cycle (ESC) testing cycle, in agreement with the Directive 1999/96/EC of the European Parliament and the Directive of the December 13<sup>th</sup>, 1999 Council,<sup>17</sup> which establishes engine and dynamometer settings, and also NO<sub>x</sub> and other pollutants emission limits. The dynamometer used in this study has a power output of 440 kW at 6000 rpm and a torque of 2334 Nm. The engine employed is in accordance with the Euro V standards, using an urea-SCR after-treatment system. Table 1 specifies the engine details.

Table 1. Engine specifications, BR- model 2012.

| Specifications         |                                     |
|------------------------|-------------------------------------|
| Emission               | Euro V "Heavy Duty"/Proconve P7     |
| Configuration          | 4 cylinders, inline                 |
| Displacement           | 4,8 liters                          |
| Bore x Stroke          | 105 x 137 mm                        |
| Combustion System      | Direct injection                    |
| Injection System       | Common Rail Electronic              |
| Aspiration             | TGV Intercooler                     |
| Power Output           | 187hp (139,7kW)<br>2,200rpm         |
| Peak Torque            | 720Nm (73kgf.m)<br>1,200 ~ 1,600rpm |
| Weight (dry)           | 426 kg                              |
| Aftertreatment         | SCR                                 |
| Dimensions (H x L x W) | 900 x 975 x 826 mm                  |

The emission data were sampled in the laboratory of vehicular emissions of the Federal University of Parana –Curitiba/Brazil, employing an engine dynamometer driving cycle using LSD (Low Sulphur Diesel - 50 ppm sulphur content), ULSD (Ultra Low Sulphur Diesel - 10 ppm sulphur content) and B20 (soybean biodiesel blended (20%) with ULSD). The main difference between LSD and ULSD is their sulphur content, which may affect SO<sub>2</sub> and particulate emissions. However, the cetane number also differs and is considered a key fuel property comprising NMHC and CO emissions.<sup>9,18</sup>

Table 2 shows the quality parameters of the reference diesel fuels and the biodiesel blend used in this research. The Standard Test Methods established by ASTM were followed. The main properties having an influence on exhaust emissions are sulphur content and cetane number, as will be discussed in the results section.

Table 2. Fuel Properties of LSD and ULSD diesel and B20 biodiesel.

| Property                                   | LSD   | ULSD  | B20   |
|--|-------|-------|-------|
| Sulphur, mg/kg                             | 24    | 4     | 6     |
| Cetane number                              | 49.2  | 53.8  | 51    |
| Glow point (°C)                            | 58.5  | 44.5  | 70.5  |
| Viscosity at 40°C (mm <sup>2</sup> /s)     | 2.6   | 3.0   | 3.15  |
| Specific mass at 20°C (kg/m <sup>3</sup> ) | 835.2 | 830.5 | 848.1 |

The gas emission data were obtained by a SESAM i60 FT, a Fourier Transform InfraRed (FTIR) multi-component measurement system from AVL. Table 3 presents some important technical characteristics of the FTIR analysis. The FTIR was calibrated to detect specific hydrocarbons (HC), nitrogen compounds (NO, NO<sub>2</sub>, N<sub>2</sub>O and NH<sub>3</sub>) and other pollutants. It also calculates NO<sub>x</sub>, total (HCD) and non-methane hydrocarbons (NMHC) concentrations. The HCD is the sum of all hydrocarbons that FTIR can analyse using a method for diesel fuel (HCD = CH<sub>4</sub>, C<sub>2</sub>H<sub>2</sub>, C<sub>2</sub>H<sub>4</sub>, C<sub>2</sub>H<sub>6</sub>, C<sub>3</sub>H<sub>6</sub>, C<sub>3</sub>H<sub>8</sub>, C<sub>4</sub>H<sub>6</sub>, nC<sub>8</sub> and AHC-aromatic hydrocarbons). The HCD expresses the total hydrocarbons (HC) for diesel emission analysis. The NMHC comprises the HCD concentration, except for the methane fraction.

Table 3. FTIR settings.

| FTIR Spectrometer Data |   |
|------------------------|---|
| Sampling Rate          | 1 scan per second (1 Hz)                      |
| Data Rate              | All measured gas components at 1 Hz           |
| Spectral Resolution    | 0.5 cm <sup>-1</sup>                          |
| Measurement Cell       | Gas cell heated to 191 °C (375.8 °F)          |
| Response Time          | t <sub>10</sub> to t <sub>90</sub> within 1 s |
| Sample Flow Rate       | 10 l/min per stream                           |
| Detector Cooling       | Liquid nitrogen, 50 ml/h                      |
| Zero/Purge Gas         | Nitrogen / Synthetic Air, 0.6 – 1.5 l/min     |
| Compressed Air         | 5 – 6 bar and max. 100 l/min per FTIR stream  |

### 3. Results and discussions

#### 3.1 Nitrogen Compounds

Analysis of Variance (ANOVA), normal probability plot of residuals and Bartlett's test of homogeneity of variances were applied to the studied compounds. The statistical analysis were performed using R software.<sup>19</sup> A preliminary analysis showed that the residuals have a normal distribution and a parametric behaviour. The Bartlett's test presented, for almost all samples, p-values less than the significance level of 0.05, confirming the homogeneity of sample variances. In conclusion, the analysis of variance results are valid, except for C<sub>2</sub>H<sub>2</sub> and C<sub>2</sub>H<sub>6</sub>.

According to the analysis of variance results the means differ due to fuel and after-treatment system choice. To analyse the interactions between fuel and after-treatment system, we applied the Tukey significant difference test. Differences between mean values at a level of  $p < 0.05$  (95% confidence level) were considered statistically significant.<sup>20</sup>

Our results, presented in Table 4, have shown that, for all studied fuels the use of the SCR system presented statistically significant different means of nitrogen oxides (NO<sub>x</sub>), nitrogen monoxide (NO) and nitrogen dioxide (NO<sub>2</sub>) emissions, compared to results when the SCR system was not used. Quantitatively, the use of the SCR system decreased NO<sub>x</sub>, NO and NO<sub>2</sub> concentrations.

According to Chin et al.<sup>1</sup>, some biodiesel blends may reduce emissions of regulated pollutants, such as PM, CO, NMHC and CO<sub>2</sub>. However, it usually increases fuel consumption and NO<sub>x</sub> emissions.

Only NO<sub>2</sub> emission means showed statistically significant differences between LSD and ULSD fuels when the engine was not equipped with the SCR system. However this trend was not observed between the ULSD and the B20 fuels. In contrast, the use of different fuels statistically affected NO<sub>x</sub>, NO and NO<sub>2</sub> emission means when the engine was equipped with the SCR system, where the highest emissions were observed for the ULSD and B20 fuels.

According to Chin et al.<sup>1</sup> and Agarwal and Das<sup>21</sup>, a NO<sub>x</sub> emission increase due to biodiesel blend fuels use, is a result of some fuel properties, such as viscosity, and also is a result of the advance in injection timing, temperature rise and abundance of oxygen available in the combustion chamber.<sup>1,21</sup> Viscosity interfere in the fuel nebulization

generating different sizes of droplets in the combustion chamber. The burning efficiency is higher with small droplets, due to a lower viscosity, leading a lower NO<sub>x</sub> emission.

Despite the fact that the WHO<sup>22</sup> has reported that sulphur content of fuels can increase NO<sub>x</sub> emissions, as it reduces catalyst efficiency, our results showed similar concentrations to all tested fuels (scenarios without SCR system), although higher concentrations using ULSD in comparison to LSD with the use of SCR system were observed.

Table 4. Average and standard deviation of exhaust emissions for nitrogen compounds (g/kWh) using SCR system on and off.

| Pollutant        | Low Sulfur Diesel |                 | Ultra Low Sulfur Diesel |               | Biodiesel B20   |               |
|------------------|-------------------|-----------------|-------------------------|---------------|-----------------|---------------|
|                  | SCR off (±SD)     | SCR on (±SD)    | SCR off (±SD)           | SCR on (±SD)  | SCR off (±SD)   | SCR on (±SD)  |
| NO <sub>x</sub>  | 7.55 ± 0.04       | 0.52 ± 0.02     | 7.66 ± 0.07             | 2.4 ± 0.8     | 7.6 ± 0.2       | 1.6 ± 0.4     |
| NO               | 4.89 ± 0.02       | 0.34 ± 0.01     | 4.84 ± 0.03             | 1.5 ± 0.5     | 4.8 ± 0.1       | 0.98 ± 0.24   |
| NO <sub>2</sub>  | 0.06 ± 0.01       | < M.D.C.        | 0.26 ± 0.04             | 0.15 ± 0.04   | 0.31 ± 0.07     | 0.06 ± 0.01   |
| NH <sub>3</sub>  | 0.004 ± 0.002     | 0.07 ± 0.02     | 0.002 ± 0.001           | 0.007 ± 0.003 | 0.0008 ± 0.0007 | 0.006 ± 0.001 |
| N <sub>2</sub> O | 0.0133 ± 0.0001   | 0.0434 ± 0.0003 | 0.0127 ± 0.0005         | 0.044 ± 0.004 | 0.013 ± 0.001   | 0.061 ± 0.008 |

NO<sub>x</sub> - Nitrogen Oxides, NO- Nitrogen Monoxide, NO<sub>2</sub>- Nitrogen Dioxide, NH<sub>3</sub>- Ammonia, N<sub>2</sub>O- Nitrous Oxide.

\* MDC (Minimal Detectable Concentration) is the detection limit of each gas component, determined as two times the standard deviation  $\sigma$  of zero gas measurement over 60 seconds.  
Inferior to MDC: NO<sub>2</sub> – Nitrogen dioxide (MDC = 0,011 g/kWh).

While designed to reduce NO<sub>x</sub> emissions, the SCR system may increase other pollutants' emissions. As demonstrated in our study, the SCR system satisfies its purpose of reducing NO<sub>x</sub> emissions. However, it brings forth new problems, such as higher emissions of N<sub>2</sub>O, NH<sub>3</sub> and some hydrocarbons.

Table 4 shows an increase in ammonia emissions due to SCR system use. The only increase considered statistically significant ( $p < 0.05$ ) was for LSD.

On the other hand, while the engine was equipped with the SCR system, there is a statistically significant difference between NH<sub>3</sub> emission means from LSD to B20 and from LSD to ULSD. The NH<sub>3</sub> emission means for ULSD and B20 could not be considered significantly different at a 95% confidence level.

Koebel et al.<sup>6</sup> reported that the SCR system uses continuous urea injections (ammonia content) to neutralize NO<sub>x</sub> emissions, which may lead to an excess of urea,



called ammonia slip. It is therefore not unreasonable to assume that the ammonia slip may be responsible for the higher  $\text{NH}_3$  emissions observed.

When the injected urea solution fails to be completely decomposed below  $200^\circ\text{C}$ , it can produce ammonium nitrate ( $\text{NH}_4\text{NO}_3$ ), cyanuric acid ( $(\text{HNCO})_3$ ), and other compounds as sub-products.<sup>8</sup> As a consequence, ammonia and ammonium salts have a relevant impact on the ecosystem, accounting for the modification of the atmosphere global radioactive balance, the reduction of atmospheric visibility, the acidification and eutrophication of the environment.<sup>23</sup>

As has been reported by European Environment Agency<sup>24</sup>, road transport contributes only 2% of total ammonia ( $\text{NH}_3$ ) emissions, though it is a significant source from a local perspective in urban areas. Many studies<sup>25-29</sup> reported that an increase in  $\text{NH}_3$  emission has occurred due to introduction of vehicles equipped with catalytic converters and adoption of urea-SCR system.

The main source of anthropogenic  $\text{N}_2\text{O}$  is agriculture,<sup>30</sup> but some concern has arisen due to new diesel exhaust after-treatment systems being responsible for  $\text{N}_2\text{O}$  production, for example, the chemical reactions in urea-SCR system.<sup>31</sup>

In our experiment, the use of the SCR system increased  $\text{N}_2\text{O}$  concentrations for all studied fuels. With 95% confidence level, these increases can be considered statistically significant, with the highest increase observed for the B20 biodiesel blend (about 361%) and the lowest for the ULSD (about 83%). These results can be explained by the undesirable processes that may occur in the SCR systems, including several competitive, non selective reactions with oxygen that can produce secondary emission.<sup>31</sup>

While the engine was equipped with the SCR system, a statistically significant increase of  $\text{N}_2\text{O}$  emission due to B20 biodiesel use was verified, in comparison with ULSD and LSD fuels ( $p < 0.05$ ).

### 3.2 Hydrocarbons

The FTIR equipment is also able to detect the non-methane hydrocarbons (NMHC) and hydrocarbons of diesel (HCD). The results are shown in Table 5.

Table 5. Average exhaust emissions for hydrocarbons compounds (g/kWh).

| Low Sulfur Diesel | Ultra Low Sulfur Diesel | Biodiesel B20 |
|-------------------|-------------------------|---------------|
|-------------------|-------------------------|---------------|

| Pollutant                     | SCR off ( $\pm$ SD) | SCR on ( $\pm$ SD)    | SCR off ( $\pm$ SD) | SCR on ( $\pm$ SD)  | SCR off ( $\pm$ SD)   | SCR on ( $\pm$ SD)  |
|-------------------------------|---------------------|-----------------------|---------------------|---------------------|-----------------------|---------------------|
| NMHC                          | 0.1888 $\pm$ 0.0002 | 0.1857 $\pm$ 0.0004   | 0.135 $\pm$ 0.003   | 0.159 $\pm$ 0.003   | 0.136 $\pm$ 0.007     | 0.164 $\pm$ 0.006   |
| HCD                           | 0.1917 $\pm$ 0.0004 | 0.1878 $\pm$ 0.0004   | 0.137 $\pm$ 0.003   | 0.161 $\pm$ 0.003   | 0.137 $\pm$ 0.007     | 0.166 $\pm$ 0.006   |
| C <sub>3</sub> H <sub>6</sub> | 0.0233 $\pm$ 0.0009 | 0.0236 $\pm$ 0.0002   | 0.012 $\pm$ 0.002   | 0.006 $\pm$ 0.001   | 0.0138 $\pm$ 0.0004   | 0.013 $\pm$ 0.003   |
| C <sub>2</sub> H <sub>2</sub> | 0.0142 $\pm$ 0.0003 | 0.0120 $\pm$ 0.0003   | 0.0125 $\pm$ 0.0008 | 0.0122 $\pm$ 0.0004 | 0.0104 $\pm$ 0.0006   | 0.0124 $\pm$ 0.0008 |
| C <sub>2</sub> H <sub>6</sub> | 0.0653 $\pm$ 0.0006 | 0.0673 $\pm$ 0.0007   | 0.064 $\pm$ 0.002   | 0.089 $\pm$ 0.003   | 0.068 $\pm$ 0.004     | 0.087 $\pm$ 0.002   |
| C <sub>3</sub> H <sub>8</sub> | 0.030 $\pm$ 0.001   | 0.0169 $\pm$ 0.0007   | 0.0276 $\pm$ 0.002  | 0.0281 $\pm$ 0.0008 | 0.0168 $\pm$ 0.0007   | 0.025 $\pm$ 0.005   |
| CH <sub>4</sub>               | 0.0028 $\pm$ 0.0003 | 0.00213 $\pm$ 0.00003 | 0.0021 $\pm$ 0.0002 | 0.0023 $\pm$ 0.0001 | 0.00165 $\pm$ 0.00007 | 0.0022 $\pm$ 0.0004 |
| HCHO                          | 0.0285 $\pm$ 0.0007 | 0.0063 $\pm$ 0.0005   | 0.011 $\pm$ 0.002   | 0.0037 $\pm$ 0.0002 | 0.010 $\pm$ 0.004     | 0.006 $\pm$ 0.002   |
| nC <sub>8</sub>               | 0.056 $\pm$ 0.001   | 0.0659 $\pm$ 0.0002   | 0.0204 $\pm$ 0.0005 | 0.024 $\pm$ 0.002   | 0.027 $\pm$ 0.002     | 0.027 $\pm$ 0.004   |

NMHC- Non-Methane Hydrocarbons, HCD- Hydrocarbons of Diesel, C<sub>3</sub>H<sub>6</sub>-Propylene, C<sub>2</sub>H<sub>2</sub>- Acetylene, C<sub>2</sub>H<sub>6</sub>- Ethane, C<sub>3</sub>H<sub>8</sub>-Propane, CH<sub>4</sub> - Methane, HCHO- Formaldehyde and nC<sub>8</sub>- N-Octane.

Inferior to MDC: C<sub>2</sub>H<sub>4</sub>- Ethene (MDC = 0,0173 g/kWh), C<sub>4</sub>H<sub>6</sub>- 1, 3 Butadiene (MDC = 0,0666 g/kWh) and AHC- Aromatic hydrocarbon (MDC = 0,0134 g/kWh).

The NMHC emission means were statistically different between LSD and ULSD for both situations, SCR-on and SCR-off, showing a reduction of 30% for SCR off and 15% for SCR on. The influence of the SCR system in NMHC emissions means was statistically significant only for ULSD and B20. The means increased by nearly 20% using ULSD and B20 ( $p < 0.05$ ). Diesel hydrocarbons emissions (HCD) showed a similar trend to that observed for NMHC emissions described previously.

Fuels with a smaller cetane number has a higher ignition delay time, which “along with the combustion of a partially premixed charge results in excessive emissions from incomplete combustion, specifically total hydrocarbons (THC) and CO”.<sup>18</sup>

Regarding recent changes on fuel properties, such as lower sulphur content in diesel and the use of biodiesel blends, considering measures of each hydrocarbon to engine not equipped with SCR system, the use of ULSD showed statistically significant difference on means in comparison to LSD to all hydrocarbons, with exception of ethane and acetylene (analysis of variance invalid). However, the only hydrocarbons showing significant differences on means ( $p < 0.05$ ) from ULSD to B20 were propane and n-octane, with decrease of propane and increase of n-octane.

Statistical treatment of data indicates that formaldehyde emissions were significantly ( $p < 0.05$ ) lower (78%) with LSD and (59%) with ULSD due to SCR system use. It also indicates that n-octane emissions were significantly ( $p < 0.05$ ) higher (18%) with LSD due to SCR system use.

Besides the toxicity of some organic compounds like BTEX and HPA's, well known as potential carcinogenic compounds, Atkinson<sup>32</sup> pointed out that a variety of hydrocarbons may lead to ozone production in low latitudes, through their reaction to OH radicals in the presence of NO<sub>x</sub> and SO<sub>2</sub>.

The ground-level ozone is a well-known atmospheric pollutant, which can cause several deleterious impacts on the environment and human health. In high concentrations, the tropospheric O<sub>3</sub> can interfere with photosynthesis and the growing of some plant species.<sup>33,22</sup> The latest European directive 2002/3/CE recommends that at least 30 NMHCs (saturated, unsaturated or aromatic) should be measured.<sup>34</sup> As far as ozone formation due to high NMHCs and SO<sub>x</sub> emissions are concerned, the critical situation in our study was that of LSD, which presented elevated NMHC and SO<sub>2</sub> emissions.

In this context, it is widely known that organic compounds participated in the formation of secondary pollutants that may contribute to some of the undesirable environmental effects associated with photochemical smog episodes.

Essentially, each compound has a different contribution due to the amount emitted and some properties that affect the secondary pollutants production during photochemical reactions. Some of these compounds are said to be more reactive than others. Consequently, the most reactive organic compounds should be addressed towards a strategy to reduce ozone and PAN (Peroxyacetylnitrate) exposure levels.<sup>35</sup>

A ranking of most reactive organic compounds, based on ozone formation under specific atmospheric conditions has been developed, the so-called reactivity scale. Derwent et al.<sup>35</sup> created a reactivity scale for Northwestern Europe. They estimated the Photochemical Ozone Creation Potentials (POCPs) and Photochemical PAN Creation Potentials (PPCPs) for 120 organic compounds and their sensitivity to NO<sub>x</sub> emissions taking ethylene (POCP = 100) and propylene (PPCP = 100), respectively, as the reference compound. Table 6 presents the values calculated by Derwent et al. (1998).<sup>35</sup>

Table 6. Photochemical Ozone Creation Potential POCP and Photochemical PAN Creation Potential

| Organic Compounds | POCP  | PPCP |
|-------------------|-------|------|
| Propylene         | 112.3 | 100  |
| Formaldehyde      | 51.9  | 14.8 |
| N-octane          | 45.3  | 42.9 |

|           |      |      |
|-----------|------|------|
| Propane   | 17.6 | 13.7 |
| Ethane    | 12.3 | 17.3 |
| Acetylene | 8.5  | 2.2  |
| Methane   | 0.6  | 0.9  |

Source: Derwent et al.<sup>35</sup>

Relating the results of Table 6 with our study, n-octane POCP is only 13% lower than formaldehyde's one, while its PPCP is 65% higher than the formaldehyde one. With regards to ozone and PAN formation, LSD fuel presented the higher concentrations for the compounds with the higher POCP and PPCP values: propylene, formaldehyde and n-octane.

Considering only the LSD fuel, it was statistically verified ( $p < 0.05$ ) an increase in n-octane emission and a decrease in formaldehyde when the SCR system was used. These results indicate a beneficial effect in ozone photochemical creation, as the formaldehyde POCP is higher than n-octane one. In addition, as reported by WHO<sup>22</sup>, formaldehyde was classified as a carcinogenic compound.

The SCR system combined with ULSD or B20 has increased alkanes emissions, however their POCP and PPCP are lower than those of formaldehyde, propylene and n-octane. Therefore, the ULSD and B20 fuels are, apparently, a better alternative than LSD, considering the hydrocarbons emissions and their photochemical potentials.

Recently Derwent et al.<sup>36</sup> developed a similar study applying the same models to create an activity scale for different emission sources of organic compounds. They indicated road transport-exhaust as the major contributor to POCP levels. Furthermore, Derwent et al.<sup>37</sup> made the same conclusion for secondary organic aerosol formation from organic compounds.

The POCP and PPCP analysis applied in our study is interesting since the combination of megacities, atmospheric conditions and significant emissions of ozone and PAN precursors can favour photochemical reactions in smog systems, creating serious pollution episodes.

Regarding the use of the SCR system scenarios, the results are of similar magnitude for all tested fuels. However, when the engine was not equipped with the SCR system, the LSD showed higher emissions, with differences over 60% in comparison to ULSD, with little difference between ULSD and B20.

Open literature describes decreases in aldehyde emissions from some biodiesel fuels, in comparison to diesel.<sup>38-40</sup> However, specifically with regard to formaldehyde, some researchers observed an increase or no alteration in its emission.<sup>41-43,9</sup> Tan et al.<sup>44</sup> showed an increase of formaldehyde emissions mainly for pure biodiesel fuel in comparison to diesel, and showed little difference between diesel and B20 blend.

Taken together, this study showed that the emissions of NO and NO<sub>2</sub> while the engine was equipped with the SCR system using the ESC cycle were lower and statistically significant ( $p < 0.05$ ). However, the use of the SCR system produced significantly increased concentrations of: N<sub>2</sub>O for all studied fuels; NH<sub>3</sub> just for LSD; and non-methane hydrocarbons (NMHC) and hydrocarbons of diesel (HCD) for ULSD and B20. On the other hand, the use of SCR system significantly ( $p < 0.05$ ) suppressed formaldehyde emissions for LSD and ULSD fuels, having a beneficial impact since it has a huge POCP and PPCP and is considered as a carcinogenic compound.

Soybean biodiesel blend used, in combination with the SCR system, can successfully reduce harmful pollutant emissions such as NO<sub>x</sub>, however, increases the HCD production.

#### 4. Acknowledgement

This work was supported financially by the National Council for Scientific and Technological Development (CNPq).

#### 5. References

- (1) Chin, J.; Batterman, S. A.; Northrop, W. F.; Bohac, S. V.; Assanis, D. N. Gaseous and Particulate Emissions from Diesel Engines at Idle and under Load: Comparison of Biodiesel Blend and Ultralow Sulfur Diesel Fuels. *Energy & Fuels*. **2012**, 26, 6737–6748; DOI 10.1021/ef300421h.
- (2) U.S. EPA. United States Environmental Protection Agency Technical Bulletin: Nitrogen oxides (NO<sub>x</sub>) why and how they are controlled. Clean Air Technology Center (MD-12), 57pp, **1999**.
- (3) Alkemade, U.; Schumann, B. Engines and exhaust after treatment systems for future automotive applications. *Solid State Ionics*. **2006**, 177, 2291–2296; DOI 10.1016/j.ssi.2006.05.051
- (4) Furfori, S.; Russo, N.; Fino, D.; Saracco, G.; Specchia, V. NO SCR reduction by hydrogen generated in line on perovskite-type catalysts for automotive diesel exhaust gas

- treatment. *Chemical Engineering Science*. **2010**, 65, 120–127; DOI 10.1016/j.ces.2009.01.065.
- (5) Miquel, P.; Granger, P.; Jagtap, N.; Umbarkar, S.; Dongare, M.; Dujardin, C. NO reduction under diesel exhaust conditions over Au/Al<sub>2</sub>O<sub>3</sub> prepared by deposition-precipitation method. *Journal of Molecular Catalysis A: Chemical*. **2010**, 322, 90–97; DOI 10.1016/j.molcata.2010.02.024.
- (6) Koebel, M.; Elsener, M.; Kleemann, M. Urea-SCR: a promising technique to reduce NO<sub>x</sub> emissions from automotive diesel engines. *Catalysis Today*. **2000**, 59, 335–345; DOI 10.1016/S0920-5861(00)00299-6.
- (7) Hu, Y.; Griffiths, K.; Norton, P.R. Surface science studies of selective catalytic reduction of NO: Progress in the last ten years. *Surface Science*. **2009**, 603, 1740–1750; DOI 10.1016/j.susc.2008.09.051.
- (8) Jiang, L.; Ge, Y.; Shah, A. N.; He, C.; Liu, Z. Unregulated emissions from a diesel engine equipped with vanadium-based urea-SCR catalyst. *Journal of Environmental Sciences*. **2010**, 22, 575–581; DOI 10.1016/S1001-0742(09)60148-0.
- (9) Turrio-Baldassarri, L.; Battistelli, C. L.; Conti, L.; Crebelli, R.; De Berardis, B.; Iamiceli, A. L.; Gambino, M.; Iannaccone, S. Emission comparison of urban bus engine fueled with diesel oil and “biodiesel” blend. *The Science of the Total Environment*. **2004**, 327, 147–62; DOI 10.1016/j.scitotenv.2003.10.033.
- (10) Lobo, F. A.; Goveia, D.; Oliveira, A. P.; Romão, L. P. C.; Fraceto, L. F.; Dias Filho, N. L.; Rosa, A. H. Development of a method to determine Ni and Cd in biodiesel by graphite furnace atomic absorption spectrometry. *Fuel*. **2011**, 90, 142–146; DOI 10.1016/j.fuel.2010.09.009.
- (11) Karavalakis, G.; Bakeas, E.; Stournas, S. Influence of oxidized biodiesel blends on regulated and unregulated emissions from a diesel passenger car. *Environmental Science and Technology*. **2010**, 44, 5306–5312; DOI 10.1021/es100831j.
- (12) Bakeas, E.; Karavalakis, G.; Stournas, S. Biodiesel emissions profile in modern diesel vehicles. Part 1: Effect of biodiesel origin on the criteria emissions. *The Science of the Total Environment*. **2011**, 409, 1670–1676; DOI 10.1016/j.scitotenv.2011.01.024.
- (13) Bermúdez, V.; Lujan, J. M.; Pla, B.; Linares, W. G. Comparative study of regulated and unregulated gaseous emissions during NEDC in a light-duty diesel engine fuelled with Fischer Tropsch and biodiesel fuels. *Biomass and Bioenergy*. **2011**, 35, 789–798; DOI 10.1016/j.biombioe.2010.10.034.
- (14) Demirbas, A. Biodiesel from oilgae, biofixation of carbon dioxide by microalgae: A solution to pollution problems. *Applied Energy*. **2011**, 88, 3541–3547; DOI 10.1016/j.apenergy.2010.12.050.
- (15) U.S. EPA. United States Environmental Protection Agency, 2013. Website <http://www.epa.gov/otaq/standards/heavy-duty/hdci-exhaust.htm>.
- (16) Koebel, M.; Madia, G.; Elsener, M. Selective catalytic reduction of NO and NO<sub>2</sub> at low temperatures. *Catalysis Today*. **2002**, 73, 239–247; DOI 10.1016/S0920-5861(02)00006-8.
- (17) EC (European Commission). European Commission Report from the Commission to the European Parliament and the Council: Quality of petrol and diesel fuel used for road transport in the European Union: Sixth annual report (Reporting year 2007), 15pp.
- (18) Lilik, G. K.; Boehman, L. Advanced Diesel Combustion of a High Cetane Number Fuel with Low Hydrocarbon and Carbon Monoxide Emissions. *Energy & Fuels*. **2011**, 25, 1444–1456; DOI 10.1021/ef101653h.
- (19) R Core Team. 2014. R: A language and environment for statistical computing. R Foundation for Statistical Computing, Vienna, Austria. Website <http://www.R-project.org/>.
- (20) Montgomery, D. C. Design and Analysis of Experiments. 5<sup>th</sup> Edition. John Wiley & Sons, INC. 699 pp, 2011.
- (21) Agarwal, A. K.; Das, L. M. Biodiesel development and characterization for use as a fuel in compression ignition engines. *Journal of Engineering for Gas Turbines and Power*. **2001**, 123, 440–447; DOI 10.1115/1.1364522.

- (22) WHO. World Health Organization. Health effects of transport-related air pollution. Copenhagen: WHO Regional Office for Europe, 205p, 2005.
- (23) Reche, C.; Viana, M.; Pandolfi, M.; Alastuey, A.; Moreno, T.; Amato, F.; Ripoll, A.; Querol, X. Urban NH<sub>3</sub> levels and sources in a Mediterranean environment. *Atmospheric Environment*. **2012**, 57, 153–164; DOI 10.1016/j.atmosenv.2012.04.021.
- (24) EEA (European Environment Agency). European Union Emission Inventory Report 1990 e 2009 under the UNECE convention on long-range transboundary air pollution. EEA Technical Report 9, 2011. Website <http://www.eea.europa.eu/publications/eu-emission-inventory-report-lrtap>.
- (25) Sutton, M. A.; Dragosits, U.; Tang, Y. S.; Fowler, D. Ammonia emissions from non-agricultural sources in the UK. *Atmospheric Environment*. **2000**, 34, 855–869; DOI 10.1016/S1352-2310(99)00362-3.
- (26) Perrino, C.; Catrambone, M.; Menno, A. Di; Bucchianico, D.; Allegrini, I. Gaseous ammonia in the urban area of Rome, Italy and its relationship with traffic emissions. *Atmospheric Environment*. **2002**, 36, 5385–5394; DOI 10.1016/S1352-2310(02)00469-7.
- (27) Cape, J. N.; Tang, Y. S.; van Dijk, N.; Love, L.; Sutton, M.; Palmer, S. C. F. Concentrations of ammonia and nitrogen dioxide at roadside verges, and their contribution to nitrogen deposition. *Environmental Pollution*. **2004**, 132, 469–478; DOI 10.1016/j.envpol.2004.05.009.
- (28) Burgard, D. A.; Bishop, G.; Stedman, D. H. Remote sensing of ammonia and sulfur dioxide from on-road light duty vehicles. *Environmental Science & Technology*. **2006**, 40, 7018–22; DOI 10.1021/es061161r.
- (29) Kean, A. J.; Littlejohn, D.; Ban-Weiss, G. A.; Harley, R. A.; Kirchstetter, T. W.; Lunden, M. M. Trends in on-road vehicle emissions of ammonia. *Atmospheric Environment*. **2009**, 43, 1565–1570; DOI 10.1016/j.atmosenv.2008.09.085.
- (30) Wang, Z.; Zheng, H.; Luo, Y.; Deng, X.; Herbert, S.; Xing, B. Characterization and influence of biochars on nitrous oxide emission from agricultural soil. *Environmental Pollution*. **2013**, 174, 289–296; DOI 10.1016/j.envpol.2012.12.003.
- (31) Majewski, W. A. 2005. Selective catalytic reduction. Website <[http://www.dieselnit.com/tech/cat\\_scr.php](http://www.dieselnit.com/tech/cat_scr.php)>.
- (32) Atkinson, R. Atmospheric chemistry of VOCs and NO<sub>x</sub>. *Atmospheric Environment*. **2000**, 34, 2063–2101; DOI 10.1016/S1352-2310(99)00460-4.
- (33) Susaya, J.; Kim, K.; Shon, Z. Demonstration of long-term increases in tropospheric O<sub>3</sub> levels: Causes and potential impacts. *Atmospheric Environment*. **2013**, 92 (11), 1520–1528; DOI 10.1016/j.chemosphere.2013.04.01.
- (34) Arsene, C.; Bougiatioti, A.; Mihalopoulos, N. Sources and variability of non-methane hydrocarbons in the eastern Mediterranean. *Global Nest Journal*. **2009**, 11 (3), 333–340; DOI 10.1007/978-1-4020-6429-6\_14.
- (35) Derwent, R. G.; Jenkin, M. E.; Saunders, S. M.; Pilling, M. J. Photochemical ozone creation potentials for organic compounds in northwest Europe calculated with a master chemical mechanism. *Atmospheric Environment*. **1998**, 32, 2429–2441; DOI 10.1016/S1352-2310(98)00053-3.
- (36) Derwent, R. G.; Jenkin, M. E.; Passant, N. R.; Pilling, M. J. Photochemical ozone creation potentials (POCPs) for different emission sources of organic compounds under European conditions estimated with a Master Chemical Mechanism. *Atmospheric Environment*. **2007**, 41, 2570–2579; DOI 10.1016/j.atmosenv.2006.11.019.
- (37) Derwent, R. G.; Jenkin, M. E.; Utembe, S. R.; Shallcross, D. E.; Murrells, T. P.; Passant, N. R. Secondary organic aerosol formation from a large number of reactive man-made organic compounds. *Science of the Total Environment*. **2010**, 408, 3374–3381; DOI 10.1016/j.scitotenv.2010.04.013.
- (38) Sharp, C. A.; Howell, S. A.; Jobe, J. Effect of Biodiesel Fuels on Transient Emissions from Modern Diesel Engines, Part II, Unregulated Emissions and Chemical Characterization. *SAE Technical*. **2000**, 01-1968, DOI: 10.4271/2000-01-1968.

- (39) Peng, C.; Yang, H.; Lan, C.; Chien, S. Effects of the biodiesel blend fuel on aldehyde emissions from diesel engine exhaust. *Atmospheric Environment*. **2008**, 42, 906–915; DOI 10.1016/j.atmosenv.2007.10.016.
- (40) Ratcliff, M. A.; Dane, A. J.; Williams, A.; Ireland, J.; Luecke, J.; McCormick, R. L.; Voorhees, K. J. Diesel Particle Filter and Fuel Effects on Heavy-Duty Diesel Engine Emissions. *Environmental Science & Technology*. **2010**, 44, 8343–8349; DOI 10.1021/es1008032.
- (41) Corkwell, K. C.; Jackson, M. M.; Daly, D. T. Review of Exhaust Emissions of Compression Ignition Engines Operating on E Diesel Fuel Blends. *SAE Technical*. **2003**, 01-3283; DOI 10.4271/2003-01-3283.
- (42) He, B. Q.; Wang, J. X.; Yan, X. G.; Tian, X.; Chen, H. Study on Combustion and Emission Characteristics of Diesel Engines Using Ethanol Blended Diesel Fuels. *SAE Technical*. **2003**, 01-0762; DOI 10.4271/2003-01-0762.
- (43) Correa, S. M.; Arbilla, G. Carbonyl emissions in diesel and biodiesel exhaust. *Atmospheric Environment*. **2008**, 42 (4), 769-775; DOI 10.1016/j.atmosenv.2007.09.073.
- (44) Tan, P. Q.; Hu, Z. Y.; Lou, D. M. FTIR detection of unregulated emissions from a diesel engine with biodiesel fuel. *Spectroscopy and Spectral Analysis*. **2012**, 32 (2), 360–363; DOI 10.3964/j.issn.1000-0593(2012)02-0360-04